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patients treated for a metastatic prostate cancer:
Cog-Pro trial protocol**

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CORRECTION

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Correction to: Impact of new generation hormone-therapy on cognitive function in elderly patients treated for a metastatic prostate cancer: Cog-Pro trial protocol

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Correction

After publication of the original article [1] the authors found that Table 2 had been formatted incorrectly, meaning that some rows in the Table did not display the correct information.

An updated version of Table 2 is included with this Correction.

The original article has also been updated.

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Table 2 Used cognitive tests, questionnaires and biological tests

Evaluations	Before inclusion	At inclusion (baseline) ^b	3 months	6 months	12 months
Signed Informed Consent	✓				
Previous medical history	✓				
Cognitive assessment^a MoCA Grober-Buschke test Digit span forward and backward (WAIS-III) Code (WAIS III) Trail Making test Doors test Stroop Victoria Verbal fluencies Rey-Osterrieth Complex Figure Number location (VOSP) Years of education and fNART		✓ Only at inclusion	✓	✓	✓
Quality of life FACT-G, FACIT-F, FACT-Cog, HADS, ISI		✓	✓	✓	✓
Pain (VAS)	✓ ^c	✓	✓	✓	✓
ONLY for PATIENTS (group of interest and control group)					
Geriatric assessment^d G8 Charlson ADL IADL MNA Time up and go		✓	✓	✓	✓
Quality of life FACT-P		✓	✓	✓	✓
Adherence evaluation^e Morisky questionnaire Patient diary			✓	✓	✓
Biological tests^f		✓	✓	✓	✓
Specific blood samples for further research^g		✓			

MoCA Montreal Cognitive Assessment, WAIS Wechsler Adult Intelligence Scale, VOSP Visual Object and Space Perception Battery, fNART French National Adult Reading Test, ISI Insomnia Severity Index, VAS Visual Analog Scale, ADL Activities of Daily Living, IADL Instrumental Activities of Daily Living, MNA Mini-Nutritional Assessment

^aCognitive assessment will be performed by neuropsychologists

^bFor group of interest patients: before the start of the treatment or within 15 days after the start of treatment by abiraterone acetate or enzalutamide

^cHad to be ≤ 3 on the 0–10 pain VAS scale to meet with inclusion pain criteria

^dGeriatric assessment will be performed by a study nurse specialized in geriatric

^eAdherence evaluation will be performed only in group of interest patients

^fAt each time: CBC, platelets, albumin, CRP, prealbumin, iron, ferritin, transferrin, creatinin, sodium, potassium, ALT, AST, GGT, ALP, total bilirubin, TSH, T4, testosterone. At inclusion only: cortisol (at 8 h AM, fasting)

^g1 EDTA (5 ml), 1 dry tube with gel (5 ml) and 1 dry tube without gel (5 ml)